# EXHIBIT CC

Paper No. 14

*All communications respecting this case should identify it by number and names of parties.*

**U.S. DEPARTMENT OF COMMERCE**
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PAT. & T.M. OFFICE
BOARD OF PATENT APPEALS
AND INTERFERENCES

Applicants: Cabilly et al
Serial No.: 07/205,419
Filed: June 10, 1988
For: RECOMBINANT IMMUNOGLOBIN PREPARATIONS
Accorded Benefit of: U.S. S.N. 06/483,457, filed 04/08/83, now Patent No. 4,816,567, issued 03/28/89

The case referred to above has been forwarded to the Board of Patent Appeals and Interferences because it is adjudged to interfere with other cases hereafter specified. Attention is directed to the fact that this interference is declared pursuant to 37 CFR 1.601 et seq., effective February 11, 1985 (49 F.R. 48416. 1050 O.G. 385). The interference is designated as No. 102,572.

By direction of the Commissioner of Patents and Trademarks and as required by 35 USC 135(c), notice is hereby given the parties of the requirement of the law for filing in the Patent and Trademark Office a copy of any agreement "in connection with or in contemplation of the termination of the interference."

The cases involved in this interference are:

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Serial No.: 06/672,265, filed 11/14/84, now Patent No. 4,816,397, issued 03/28/89

For: MULTICHAIN POLYPEPTIDES OR PROTEINS AND PROCESSES FOR THEIR PRODUCTION

Assignee: Celltech Limited, Berkshire SL1 4DY, U.K., A British Company

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Serial No.: 07/205,419, filed 06/10/88

For: RECOMBINANT IMMUNOGLOBULIN PREPARATIONS

Assignee: Genentech, Inc., South San Francisco, CA, A California Corporation

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Accorded Benefit of: U.S. S.N. 06/483,457, filed 04/08/83, now Patent No. 4,816,567, issued 03/28/89

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Count 1

A process for producing an Ig molecule or an immunologically functional Ig fragment comprising at least the variable domains of the Ig heavy and light chains, in a single host cell, comprising the steps of:

(i) transforming said single host cell with a first DNA sequence encoding at least the variable domain of the Ig heavy chain and a second DNA sequence encoding at least the variable domain of the Ig light chain, and

(ii) independently expressing said first DNA sequence and said second DNA sequence so that said Ig heavy and light chains are produced as separate molecules in said transformed single host cell.

The claims of the parties which correspond to this count are:

Boss et al: Claims 1-18.

Cabilly et al: Claims 101-120.

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MFD/raj